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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO. CONFIRMATION N		
10/511,409	10/13/2004	Xia Xu	4239-64801-02	9192	
	7590 10/23/2009 WASHBURN LLP		EXAMINER		
CIRA CENTRE 2929 ARCH ST	E, 12TH FLOOR		FRITCHMAN, REBECCA M		
	IA, PA 19104-2891		ART UNIT	PAPER NUMBER	
			1797		
		MAIL DATE	DELIVERY MODE		
			10/23/2009	PAPER	

Please find below and/or attached an Office communication concerning this application or proceeding.

The time period for reply, if any, is set in the attached communication.

Office Action Summary		Applica	tion No.	Applicant(s)		
		10/511,	409	XU ET AL.		
		Examin	er	Art Unit		
		REBEC	CA FRITCHMAN	1797		
The Period for Rep	MAILING DATE of this commun	nication appears on t	he cover sheet with the	correspondence a	ddress	
A SHORTE WHICHEVI - Extensions o after SIX (6) - If NO period - Failure to rep Any reply rec	ENED STATUTORY PERIOD F ER IS LONGER, FROM THE N f time may be available under the provision MONTHS from the mailing date of this com for reply is specified above, the maximum s ly within the set or extended period for repl eived by the Office later than three months t term adjustment. See 37 CFR 1.704(b).	MAILING DATE OF sof 37 CFR 1.136(a). In nomination. tatutory period will apply and y will, by statute, cause the a	FHIS COMMUNICATIO event, however, may a reply be ti will expire SIX (6) MONTHS fror pplication to become ABANDON	N. mely filed n the mailing date of this ED (35 U.S.C. § 133).		
Status						
2a)⊠ This 3)⊡ Since	consive to communication(s) fil action is FINAL . In this application is in condition In din accordance with the pract	2b)∏ This action is n for allowance excep	non-final. ot for formal matters, pr		e merits is	
Disposition of	Claims					
4a) O 5) ☐ Clain 6) ☑ Clain 7) ☐ Clain	n(s) <u>1-18</u> is/are pending in the f the above claim(s) is/an(s) is/an(s) is/an(s) is/are allowed. n(s) <u>1-18</u> is/are rejected. n(s) is/are objected to. n(s) are subject to restrict appers	are withdrawn from c				
		a Evaminar				
10)☐ The d Applic Repla	pecification is objected to by the rawing(s) filed on is/are cant may not request that any objectement drawing sheet(s) including ath or declaration is objected the same of the content of the con	e: a) accepted or lection to the drawing(s g the correction is requ	be held in abeyance. Se uired if the drawing(s) is ol	ee 37 CFR 1.85(a). ojected to. See 37 C		
Priority under	35 U.S.C. § 119					
 12) Acknowledgment is made of a claim for foreign priority under 35 U.S.C. § 119(a)-(d) or (f). a) All b) Some * c) None of: 1. Certified copies of the priority documents have been received. 2. Certified copies of the priority documents have been received in Application No 3. Copies of the certified copies of the priority documents have been received in this National Stage application from the International Bureau (PCT Rule 17.2(a)). * See the attached detailed Office action for a list of the certified copies not received. 						
2) 🔲 Notice of Dr	oferences Cited (PTO-892) aftsperson's Patent Drawing Review (Disclosure Statement(s) (PTO/SB/08) IMail Date		4) Interview Summar Paper No(s)/Mail [5] Notice of Informal 6) Other:	Oate		

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Detailed Action Summary

1. This is Final Office action based on 10/511409 application attorneys remarks filed on 06/22/2009.

2. Claims 1-18 are pending and have been fully considered.

Claim Rejections - 35 USC § 103

3. Claims 1-9, 12-14, & 16-18 are rejected under 35 U.S.C. 103(a) as being obvious over VISSER et al(High-performance liquid chromatography of the neuroactive steroids alphaxalone and prefnanolone in plasma using dansyl hydrazine as fluorescent label: application to pharmacokinectic-pharmacodynamic study in rats, J Chrom B, vol. 745, pgs. 357-363(2000)) in view of Bailey et al.(USP 5807748), and in further view of TAKADATE et al. in (A Convenient Derivatization with Anion Exchange Resin Catalysts for High-Performance Liquid Chromatographic Analysis. I. Derivatization of Estrogens with Dansyl Chloride, Chem. Pharm. Bull., vol. 33, no. 11, pages 5092-5095(1985)).

VISSER et al. teaches of reacting a sample with a sulfohydrazine to form a sulfonhydrazone of a ketosteroid (Figure 1) and analyzing the reacted sample by high-performance liquid chromatography (abstract), wherein detection of the sulfonhydrazone indicates presence of the ketosteroid (Section 3). The non-polar phase is a C18 stationary phase (Section2) A methanol/water solvent is used during the HPLC (Section 2).

VISSER et al. does not teach of analyzing the reacted sample by mass spectrometry. BAILEY et al. teach of using reversed phase HPLC and electrospray mass spectrometry for separating and identifying a fluorescent derivative of an analyte molecule (Columns 1-4, examples). Bailey et al. teaches that the disclosed method improves sensitivity (column 4). Therefore, it would have been obvious to one of ordinary skill in the art to modify the method of VISSER et al with the separation and identification techniques of BAILEY et al. in order to gain the advantages of improved detection sensitivity.

VISSER and BAILEY et al. do not teach of using a sulfonyl halide following using the sulfonhydrazide. TAKADATE et al. teach that derivatization of ketosteroid estrogens using a sulfonyl halide is an alternative derivatization to that of ketosteroids using sulfonhydrazides. It would have been obvious to one of ordinary skill in the analytical chemistry arts to follow the derivatization of ketosteroids using sulfonhydrazides with an alternative and well-known method of derivatization of ketosteroids using sulfonyl halide. The purpose or advantage of doing so would be to check the relative accuracy of the first method. Thus, it would have been obvious for one of ordinary skill in the art to follow the sulfonhydrazide reaction with a sulfonyl halide reaction to gain the advantages of comparison testing.

4. Claims 1-18 are rejected under 35 U.S.C. 103(a) as being obvious

over BERLINER in US 5272134 in view of Bailey in US 5807748, and in

further view of TAKADATE et al. in (A Convenient Derivatization with Anion

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Exchange Resin Catalysts for High-Performance Liquid Chromatographic

Analysis. I. Derivatization of Estrogens with Dansyl Chloride, Chem. Pharm.

Bull., vol. 33, no. 11, pages 5092-5095(1985)).

BERLINER teaches of reacting a sample with sulfonhydrazide to form a sulfonhydrazone of a ketosteroid in the sample and analyzing the reacted sample (Example 13-15). Berliner teaches that using methanol washes the product clean (Examples 13-15) and teaches using the methanol/water elution ratio 80:20(examples 13-15).

BERLINER does not expressly teach the method of analysis used.

BAILEY et al. teach of using reversed phase HPLC and electrospray mass spectrometry for separating and identifying a fluorescent derivative of an analyte molecule (Columns 1-4, examples). Bailey et al. teaches that the disclosed method improves sensitivity (column 4). Therefore, it would have been obvious to one of ordinary skill in the art to modify the method of BERLINER et al with the separation and identification techniques of BAILEY et al. in order to gain the advantages of improved detection sensitivity.

BERLINER and BAILEY et al. do not teach of using a sulfonyl halide following using the sulfonhydrazide. TAKADATE et al. teach that derivatization of ketosteroid estrogens using a sulfonyl halide is an alternative derivatization to that of ketosteroids using sulfonhydrazides. It would have been obvious to one of ordinary skill in the analytical chemistry arts to follow the derivatization of ketosteroids using sulfonhydrazides with an alternative and well-known method of derivatization of ketosteroids using sulfonyl halide. The purpose or advantage

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of doing so would be to check the relative accuracy of the first method. Thus, it would have been obvious for one of ordinary skill in the art to follow the sulfonhydrazide reaction with a sulfonyl halide reaction to gain the advantages of comparison testing.

Response to Arguments

Applicant's arguments filed 06/22/2009 have been fully considered but they are not persuasive.

In response to applicant's argument that the examiners reasoning for combination(with TAKADATE) is different than the applicants, which is that, "the two-step process of carbonyl derivatization followed by hydroxyl derivatization provides for better HPLC separation of steroids, and allows for better signal detection in API-MS", the fact that applicant has recognized another advantage which would flow naturally from following the suggestion of the prior art cannot be the basis for patentability when the differences would otherwise be obvious. See *Ex parte Obiaya*, 227 USPQ 58, 60 (Bd. Pat. App. & Inter. 1985).

Conclusion

Applicant's amendment necessitated the new ground(s) of rejection presented in this Office action. Accordingly, **THIS ACTION IS MADE FINAL**. See MPEP § 706.07(a). Applicant is reminded of the extension of time policy as set forth in 37 CFR 1.136(a).

A shortened statutory period for reply to this final action is set to expire THREE MONTHS from the mailing date of this action. In the event a first reply is

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filed within TWO MONTHS of the mailing date of this final action and the advisory action is not mailed until after the end of the THREE-MONTH shortened statutory period, then the shortened statutory period will expire on the date the advisory action is mailed, and any extension fee pursuant to 37 CFR 1.136(a) will be calculated from the mailing date of the advisory action. In no event, however, will the statutory period for reply expire later than SIX MONTHS from the date of this final action.

Any inquiry concerning this communication or earlier communications from the examiner should be directed to REBECCA FRITCHMAN whose telephone number is (571)270-5542. The examiner can normally be reached on Monday-Friday 7:30-5:00.

If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Kim, Vickie can be reached on 571-272-0579. The fax phone number for the organization where this application or proceeding is assigned is 571-273-8300.

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Information regarding the status of an application may be obtained from the Patent Application Information Retrieval (PAIR) system. Status information for published applications may be obtained from either Private PAIR or Public PAIR. Status information for unpublished applications is available through Private PAIR only. For more information about the PAIR system, see http://pair-direct.uspto.gov. Should you have questions on access to the Private PAIR system, contact the Electronic Business Center (EBC) at 866-217-9197 (toll-free). If you would like assistance from a USPTO Customer Service Representative or access to the automated information system, call 800-786-9199 (IN USA OR CANADA) or 571-272-1000.

/Krishnan S Menon/ Primary Examiner, Art Unit 1797

R.F.